

AKTIA!

Init I1

EN – User Manual
FR – Manuel d'utilisation
DE – Bedienungsanleitung
IT – Manuale d'uso



ENGLISH

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1 Introduction

Thank you for selecting Aktiia arm type blood pressure monitor (Aktiia Init I1). The monitor features include blood pressure measurement and pulse rate measurement. The monitor is designed for at least two years of reliable service.

Readings taken by the Aktiia Init I1 are equivalent to those obtained by a trained observer using the cuff and stethoscope auscultation method. This manual contains important safety and care information and provides step-by-step instructions for using the product. Please read the manual thoroughly before using the product.



Please read this User Manual carefully to gain a complete understanding of the device's functions and safety-related information.

Support: In case you have any additional questions, or you encounter any issues, please contact Aktiia's Customer Service through our Help Center / FAQ (Frequently Asked Questions) on our website at www.aktiia.com.

Incident and Adverse Event: For any incident or event, please contact immediately Aktiia at support@aktiia.com with mention of Incident or Adverse Event in the email content or title.

2 Intended purpose

Aktiia Init I1 is an oscillometric blood pressure monitor intended to initialize the Aktiia Bracelet G1 which is a component of the Aktiia Blood Pressure Monitoring System. The Aktiia Init I1 helps with measurement of baseline blood pressure and heart rate of a user to initialize the Bracelet G1 device.



Please read the User Manual of the Bracelet G1 device.

3 Indications for use

The Aktiia Init I1, referred to as the Aktiia Cuff, is used to initialize the Aktiia Bracelet G1. The Aktiia Cuff is indicated to be used for measuring blood pressure and heart rate in adults for home use only.

4 Contraindications

- The Aktiia Cuff should not be used by any person who is pregnant or may possibly be pregnant.
- The Aktiia Cuff is not suitable for use on patients with implanted electrical devices, such as cardiac pacemakers or defibrillators.
- The Aktiia Cuff is not suitable for use neonatal patients, children, patients with pre-eclampsia, premature ventricular beats, atrial fibrillation, peripheral arterial disease, and patients undergoing intravascular therapy or arterio-venous shunt, or people who have received a mastectomy.

Please consult your doctor prior to using the device if you have one of these listed conditions.

5 Intended population

The Aktiia Cuff is intended to be used by adult population with arm circumference ranging from 22cm to 42cm (about 8¾" - 16½"). There are no restrictions related to the user's sex and ethnicity.

6 Technological characteristics



This product uses the Oscillometric Measuring method to detect blood pressure. Before every measurement, the unit establishes a "zero pressure" equivalent to the atmospheric pressure. With inflation of the arm cuff, the unit detects pressure oscillations generated by the brachial artery pulsatility, which are used to determine the systolic and diastolic pressure, as well as pulse rate.

7 Important safety information


Please read the important safety information in this user manual before using the device.

Any serious incident occurring in relation to Aktiia Cuff should be reported to Aktiia and the competent authority of the Member State in which the user and / or patient is established.

Warning and Caution symbols definition:

Symbol	Definition	Symbol	Definition
	The "WARNING" sign throughout this user manual indicates a potentially hazardous situation which, if not avoided, could result in serious injury or death.		The "CAUTION" sign throughout this user manual indicates a potentially hazardous situation which, if not avoided, could result in minor injury to the user or patient or damage to the equipment or other property.

7.1 Warnings

	<ul style="list-style-type: none">• Keep the device out of reach of young children to avoid swallowing of small parts.• This device is intended for non-invasive measuring and monitoring of arterial blood pressure. It is not intended for use on extremities other than the arm or for functions other than obtaining a blood pressure measurement.• If you are taking medications, consult your physician to determine the most appropriate time to measure your blood pressure. Never change a prescribed medication without consulting your physician.• Do not apply the cuff over a wound, otherwise it can cause further injury.
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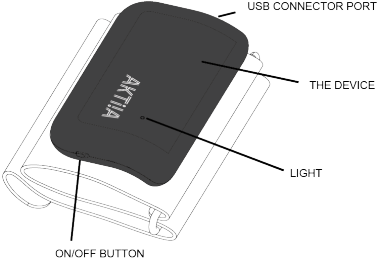


7.2 Cautions



- This device may be used only for the purpose described in this manual. The manufacturer cannot be held liable for damage caused by incorrect application.
- Do not inflate the cuff on the same limb which other monitoring equipment is applied simultaneously. This could cause temporary loss of function of the monitoring equipment that is being simultaneously used.
- On the rare occasion of a fault causing the cuff to remain fully inflated during measurement, open the cuff immediately. Prolonged high pressure (cuff pressure > 300mmHg or constant pressure < 15mmHg for more than 3 minutes) applied to the arm may lead to an ecchymosis.
- Check that operation of the device does not result in prolonged impairment of the patient's blood circulation. Too frequent and consecutive measurements could cause disturbances in blood circulation and injuries.
- This device comprises sensitive components and must be treated with caution. Observe the storage and operating conditions described in this manual.
- Do not realize servicing / maintenance while the device is in use. It is recommended that the performance should be checked every 2 years, as well as after maintenance and repair. Contact the manufacturer for such operation.
- If you are allergic to polyester, nylon or plastic, don't use this device.
- Check the device before use, do not use the device if it is damaged in any way. The use of a damaged unit may cause injury or improper results.
- The service life of the cuff may vary by the frequency of measurement and cleaning and storage state. The typical service life is 10,000 measurements.
- Please dispose the device and associated accessories according to local disposal guidelines.
- Dust may affect the performance of the unit. Please use the soft cloth to clean the whole unit before and after use. Don't use any abrasive or volatile cleaners.
- Don't open or repair the device by yourself in the event of malfunctions. The device must only be serviced, repaired and opened by individuals at authorized sales/ service centers.
- Please use accessories and detachable parts specified / authorised by the manufacturer. Otherwise, it may cause damage to the unit or danger to the user.

8 Package content

Your Aktiia Cuff is supplied in a box containing the following items:

1x Aktiia Init I1	1x USB cable	1x User Manual
 <p>Labels in the diagram:</p> <ul style="list-style-type: none">USB CONNECTOR PORTTHE DEVICELIGHTON/OFF BUTTON		 <p>Latest version of Aktiia Init I1 user manual can also be found online at: www.aktiia.com/ifu</p>

If the Aktiia packaging or any other components are damaged or open upon reception, do not use and contact Aktiia's Customer Service through our Help Center / FAQ (Frequently Asked Questions) on our website at www.aktiia.com.

9 Power supply and charging

The battery of Aktiia Cuff is a built-in rechargeable li-polymer battery. The battery capacity is 1000 mAh. If charging for the first time (immediately after purchase), after not having used it for a long time, or if the battery stops working while using the device - make sure to fully charge the Aktiia Cuff.



To charge your device:

1. **Switch ON** your Aktiia Cuff, the light indicator will flash blue.
2. Connect the USB microB connector of the provided USB cable to the device's connector port.
3. Connect the USB A connector of the USB cable to a power source with an output of 5V and 1A.
4. The light indicator will flash green to indicate charging and stay green when fully charged.

Charge the battery under following circumstances:

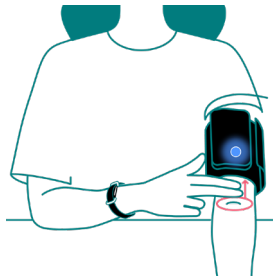
- When the red light is flashing, the battery power is low.
- When powering on the monitor, the light doesn't light up

Note: Charge at least once every three months. If the battery completely loses all charge, it may not be rechargeable anymore.

	Batteries shall not be exposed to excessive heat such as direct sunshine, fire, or other similar situations. The battery could explode causing injury or death.
	Do not attempt to replace the device battery: it is built-in and not changeable. Only charge the battery in accordance with the user instructions supplied with the device. Do not use the blood pressure monitor while charging. Do not clean the blood pressure monitor when it is being charged.

10 Correct Aktiia Cuff positioning

1. Remove garments from your upper arm. If you roll up your sleeve, please ensure that the garment is not too tight so that it does not cause any blood flow constriction.
2. Place your bare arm through the cuff to position the cuff 2-3 cm (1") above your elbow joint with the blue light indicator closest to your elbow.
3. Secure and tighten the cuff around your arm so that it fits closely but you can still insert two fingers between your arm and the cuff. Please note that if the cuff is too loose, the measurement will not be accurate.
4. While seated, place your hand, palm-side up in front of you so that it is supported by a flat surface and the Aktiia Cuff is at the same height as your heart. Your Aktiia Cuff is positioned on the inner side of your arm, over the artery.



11 Body posture during measurement

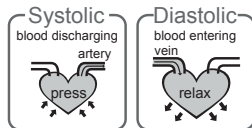
Please sit down and relax for 5 minutes before starting the initialization procedure.

1. Sit upright with your back straight and your feet flat on the floor. Do not cross your legs.
2. Place your hand palm-side down in front of you on a flat surface such as a desk or a table.
3. The middle of the Aktiia Cuff should be placed at the same level as your heart.
4. Do not move or tense your arm muscles during measurement.
5. Relax, and do not move or talk.

Note: Blood Pressure measurements can be affected by the position of the cuff and your physiological and emotional condition.

12 Important facts about Blood Pressure measurement

Blood pressure is the pressure applied by circulating blood on the walls of blood vessels. Blood pressure is mainly due to the work of the heart pumping blood through the circulatory system. When the ventricles contract and pump blood out of the heart, the blood pressure reaches its maximum value in the cardiac cycle, which is called systolic pressure. When the ventricles relax, the blood pressure reaches its minimum value in the cycle, which is called diastolic pressure.



Individual blood pressure naturally varies through regular daily life. Some circumstances have a larger impact on your blood pressure variation and may bias the measurement.



Measurements may be inaccurate if taken in the following circumstances:



Within 1 hour after eating or drinking



Immediate measurement after tea, coffee, smoking



Within 20 minutes after taking a bath



When talking or moving your fingers



In a very cold environment



When you want to discharge urine

13 How to evaluate your Blood Pressure



The Aktiia App and devices are not intended to be a diagnostic device. Self-diagnosis of measurement results and self-treatment are potentially dangerous. You should always consult your doctor for relevant interpretation and diagnosis based on your personal blood pressure results.

The following classifications are based on measurements taken on a seated person after few minutes of rest.

It is important to note that Blood Pressure readings in normal life conditions might be higher.

The table is not intended to provide a basis for any type of diagnosis or emergency assessment; the table only depict different classifications of blood pressure.

Consult your physician for an interpretation and diagnosis based on your personal blood pressure results.

Europe

The European Society of Hypertension (ESH) has created the following guide for classifying blood pressure values.

BLOOD PRESSURE CATEGORY	SYSTOLIC BP mmHg		DIASTOLIC BP mmHg	COLOR INDICATOR
OPTIMAL	LESS THAN 120	AND	LESS THAN 80	Dark Green
NORMAL	120-129	AND/OR	80-84	Light Green
ELEVATED	130-139	AND/OR	85-89	Yellow
HIGH BLOOD PRESSURE STAGE I	140-159	AND/OR	90-99	Orange
HIGH BLOOD PRESSURE STAGE 2	160-179	AND/OR	100-109	Light Red
HIGH BLOOD PRESSURE STAGE 3	HIGHER THAN 179	AND/OR	HIGHER THAN 109	Dark Red



Various factors such as age, obesity and medical condition should be considered for a correct evaluation. Consult with your physicians for an accurate assessment and diagnosis of your health condition.

14 Some frequently asked questions

Why does my blood pressure measurements differ throughout the day?

Individual blood pressure naturally varies through regular daily life (see §10). It is also affected by the way you tie your cuff and your measurement position, so please try take the measurements under the same conditions. If you are under prescription drugs, your blood pressure may vary more.

Why do I get a different blood pressure at home compared to the hospital?

The blood pressure is different even throughout the day due to weather, emotion, exercise, etc. Also, there is a phenomenon known as "white coat" effect; evidence shows that blood pressure usually increases in clinical settings, due to stress, anxiety, or other causes.

What you need to pay attention to when you measure your blood pressure at home?

- The cuff is tied properly.
- The cuff is not too tight or too loose.
- The cuff is tied on the upper arm.
- You are relaxed. Waiting for 5 minutes and taking deep breaths before beginning will yield a more accurate measurement.

Is the result the same if measuring on the right arm?

- It is ok to measure on either arm, but the results may vary for different people.
- We suggest you measure on the same arm every time.

15 Unpair your device

During the pairing procedure, the Aktiia Cuff is linked to your Aktiia account. Suppose you wish to allow another person to use the Aktiia Cuff. In that case, you must unpair the device from your Aktiia account.

To complete the unpairing procedure you should:

1. Login to your account
2. Tap on the device tab
3. Tap on the button with the three dots
4. Press unpair

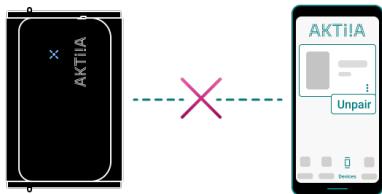
Note: Unpairing is needed if Aktiia Cuff need to be linked with a new user account.

Note: Unpairing is not needed if a new mobile device is used with the same user account.

16 Care and maintenance

To obtain the best performance, please follow the instructions below.

- Store in a dry place and avoid sunshine.
- Avoid intense shaking, or collisions.
- Use a slightly damp cloth to remove any dirt or dust.
- Avoid immersing in the water. Clean with a dry cloth if wet.
- Avoid dusty environments or fluctuating temperatures.
- Avoid washing the device other than with a damp cloth (per above).



17 Troubleshooting

Problem	Symptom	Check this	Solution
No power	The status LED is not on	The battery is empty	Recharge the device
		The side button is on the OFF position	Switch the button to the ON position
Low battery	Status LED flashing red	Battery is low	Recharge the device
Cuff not inflating	The cuff does not inflate during initialization	The cuff is switched ON and charged	Switch the cuff ON, charge fully, or contact customer support

18 Warranty

Your Aktiia Init I1 is warranted to be free from defects in materials and workmanship within two years from the date of purchase when used in accordance with the provided instructions. The warranty extends only to the end user. We will, at our option, repair or replace without charge Aktiia Init I1 covered by the warranty. Repair or replacement is our only responsibility and your only remedy under the warranty.



Do not attempt to disassemble the device as this will result in permanent damages and will void your warranty.

19 Specifications


Power supply	3.7V 1000mAh Built-in rechargeable Power supply li-polymer battery, 5V 1AAC Adaptor (Optional)
Measurement mode	Oscillometric testing mode
Measurement range	Rated cuff pressure: 0mmHg~299mmHg (0kPa ~ 39.9kPa) Measurement pressure: SYS: 60mmHg~230mmHg (8.0kPa~30.7kPa) DIA: 40mmHg~130mmHg (5.3kPa~17.3kPa) Pulse value: (40-199)beat/minute
Accuracy (according to the clinical evaluation)	Aktiia Init I1 complies with the accuracy requirements of ISO 81060-2 Blood Pressure: Mean error \pm 5mmHg / Standard deviation \pm 8mmHg Pulse value: \pm 5%
Normal working condition	A temperature range of: +5°C to +40°C A relative humidity range of 15% to 90%, non-condensing, but not requiring a water vapour partial pressure greater than 50 hPa An atmospheric pressure range of: 700 hPa to 1060 hPa
Storage & transportation condition	Temperature: -5°C to +50°C A relative humidity range of \leq 93%, non-condensing, at a water vapour pressure up to 50hPa
Measurement perimeter of the arm	About 22cm-42cm (8 $\frac{3}{4}$ to 16 $\frac{1}{2}$ inches)





Weight	Approx.271g
External dimensions	Approx.74.3mm×28.2mm×133mm
Attachment	USB Cable and user manual
Mode of operation	Continuous operation
Degree of protection	Type BF applied part
Protection against ingress of water	IP22: The first number 2: Protected against solid foreign objects of 12,5mm Φ and greater. The second number: Protected against vertically falling water drops when enclosure titled up to 15°. Vertically falling drops shall have no harmful effects when the enclosure is titled at any angle up to 15° on either side of the vertical.
Device classification	Battery Powered Mode: Internally Powered Medical Electrical Equipment AC Adaptor charged Mode: Class II Medical Electrical Equipment (the optional AC Adaptor shall comply with the requirement of IEC 60601-1 or 60950)
Data transfer	Bluetooth Low Energy (BLE) Operating Frequency: 2402 MHz – 2480 MHz Type of Modulation: GFSK Transmission power: max. 4 dBm

20 EMC and RF statements

Aktiia Init I1 needs special precautions regarding EMC and needs to be installed and put into service according to the EMC information provided in the following section.

The device is suitable for home healthcare environments.

Aktiia Init I1 is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. Interference may occur in the vicinity of equipment marked with the following symbol .

	Use of this equipment adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this equipment and the other equipment should be observed to verify that they are operating normally.
	Use of accessories, transducers and cables other than those specified or provided by the manufacturer of this equipment could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation.
	Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the device, including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.
	Aktiia Init I1 is not suitable for use in MRI (Magnetic Resonance Imaging) environment.

21 Electromagnetic compatibility information

Guidance and manufacture's declaration - electromagnetic emissions

Aktiia Init I1 is intended for use in the electromagnetic environment specified below.

The user of Aktiia Init should ensure that it is used in such an environment.

Aktiia Init I1 is suitable for use in "Home Healthcare Environment", i.e. all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.

Emission test	Compliance	Electromagnetic environment - guidance
Conducted emissions CISPR11	Groupe 1	Aktiia Init I1 uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
Radiated emissions CISPR11	Class B	
Harmonic emissions IEC 61000-3-2	Class A	—
Voltage fluctuations/Flicker emissions IEC 61000-3-3	Compliant	—

Guidance and manufacturer's declaration - electromagnetic immunity

Aktiia Init I1 is intended for use in the electromagnetic environment specified below. The user of Aktiia Init I1 should ensure that it is used in such an environment. Aktiia Init I1 is suitable for use in "Home Healthcare Environment", i.e. all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.

Immunity test	60601-1-2 test levels	Compliance level
Electrostatic discharge (ESD) IEC 61000-4-2	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV in air	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV in air
Electrical fast transient/burst IEC 61000-4-4	± 2 kV for power supply lines ± 1 kV signal input/output 100 kHz repetition frequency	± 2 kV for power supply lines ± 1 kV signal input/ output 100 kHz repetition frequency
Surge IEC61000-4-5	± 0.5 kV, ± 1 kV differential mode ± 0.5 kV, ± 1 kV, ± 2 kV common mode	± 0.5 kV, ± 1 kV differential mode ± 0.5 kV, ± 1 kV, ± 2 kV common mode
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	0 % UT; 0,5 cycle. At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315°.0 % UT; 1 cycle and 70 % UT; 25/30 cycles; Single phase: at 0°.0 % UT; 250/300 cycle	0 % UT; 0,5 cycle. At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315°.0 % UT; 1 cycle and 70 % UT; 25 / 30 cycles; Single phase : at 0°.0 % UT; 250 / 300 cycle
Power frequency magnetic field IEC 61000-4-8	30 A/m 50Hz/60Hz	30 A/ m 50Hz/ 60Hz

Conducted RF IEC61000-4-6	3 V 0,15 MHz – 80 MHz 6 V in ISM and amateur radio bands between 0,15 MHz and 80 MHz 80 % AM at 1 kHz	3 V 0,15 MHz – 80 MHz 6 V in ISM and amateur radio bands between 0,15 MHz and 80 MHz 80 % AM at 1 kHz
Radiated RF IEC61000-4-3		
Note: UT is the a.c. mains voltage prior to application of the test level.		

Guidance and manufacturer's declaration - electromagnetic immunity							
Radiated RF IEC61000-4-3	Test Frequency (MHz)	Band (MHz)	Service	Modulation	Modulation (W)	Distance (m)	Immunity Test Level (V/m)
(Test specifications for ENCLOSURE PORT IMMUNITY to RF wireless communications equipment)	385	380-390	TETRA 400	Pulse modulation b) 18Hz	1.8	0.3	27
	450	430-470	GMRS 460, FRS 460	FM c) ± 5 kHz deviation 1 kHz sine	2	0.3	28

	710	704-787	LTE Band 13,17	Pulse modulation b) 217Hz	0.2	0.3	9
	745						
	780						
	810	800-960	GSM 800/900, TETRA 800, iDEN 820, CDMA 850, LTE Band 5	Pulse modulation b) 18Hz	2	0.3	28
	870						
	930						
	1720	1700-1990	GSM 1800, CDMA 1900, GSM 1900, DECT, LTE Band 1,3,4,25, UMTS	Pulse modulation b) 217Hz	2	0.3	28
	1845						
	1970						
	2450	2400-2570	Bluetooth, WLAN, 802.11 b/g/n, RFID 2450, LTE Band 7	Pulse modulation 217Hz	2	0.3	28
	5240	5100-5800	WLAN 802.11 a/n	Pulse modulation 217Hz	0.2	0.3	9
	5500						
	5785						

22 CE compliance

This device complies with the following regulations and normative documents/standards:

EU RED STATEMENT: Hereby, Aktiia SA, declares that the device is compliance with the essential requirements and other relevant provisions of RE Directive 2014/53/EU.

EN 60601-1:2006+A1:2013/ IEC 60601-1:2005+A1:2012 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance

EN 60601-1-2:2015/ IEC 60601-1-2:2014 Medical electrical equipment – Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic disturbances - Requirements and tests

EN 60601-1-6:2010+A1:2015/IEC 60601-1-6:2010+A1:2013 Medical electrical equipment – Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability

EN 60601-1-11:2015/ IEC 60601-1-11:2015 Medical electrical equipment – Part 1-11: General requirements for basic safety and essential performance - Collateral standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment

IEC 80601-2-30:2009+A1:2013 Medical electrical equipment – Part 2-30: Particular requirements for the basic safety and essential performance of automated non-invasive sphygmomanometers

EN 1060-3:1997+A2:2009 Non-invasive sphygmomanometers – Part 3: Supplementary requirements for electro-mechanical blood pressure measuring systems

IEC 62366-1:2015 Medical devices – Part 1: Application of usability engineering to medical devices

EN 62304:2006/AC: 2008 / IEC 62304: 2006+A1:2015 Medical device software - Software life-cycle processes

ISO 10993-5:2009 Biological evaluation of medical devices – Part 5: Tests for in vitro cytotoxicity

ISO 10993-10:2010 Biological evaluation of medical devices – Part 10: Tests for irritation and skin sensitization

23 Disposal























Actuation of European directives 2002/95/EC, 2002/96/EC and 2003/108/EC, for reduction in use of dangerous substances in the electric and electronic device and for garbage disposal. The symbol applied on the device or its packaging means that at the end of its useful life the product must not be disposed of with domestic waste.

At the end of the device's useful life, the user must deliver it to the able collecting centers for electric and electronic garbage or give back to the retailer when purchasing a new device. Disposing of the product separately prevents possible negative consequences for the environment and for health, deriving from inadequate disposal. It also allows the recovery of materials of which it's made up in order to obtain an important saving of energy and resources and to avoid negative effects to the environment and health. In case of abusive disposal of device by the user, will be applied administrative endorsements in compliance with current standard. The device and its parts are made with regard to disposal, as appropriate, in accordance with national or regional regulations.

This product complies with RoHS Directive 2011/65/EU and Amendment (EU) 2015/863.

24 Safety symbols definitions

The signs below might be in the user manual, labeling or other component with your Aktia Init I1.

Symbol	Definition	Symbol	Definition
	The "WARNING" sign throughout this user manual indicates a potentially hazardous situation which, if not avoided, could result in serious injury or death.		The "CAUTION" sign throughout this user manual indicates a potentially hazardous situation which, if not avoided, could result in minor injury to the user or patient or damage to the equipment or other property.
	Symbol for "IFU are available on www.Aktia.com/ifu "		Symbol for "THE INSTRUCTION FOR USE MUST BE READ"
	Symbol for "COMPLIES WITH MDR EU2017/745 REQUIREMENTS"		Symbol for "ENVIRONMENT PROTECTION - Electrical waste products should not be disposed with household waste. Please recycle where facilities exist. Check with your local authority or retailer for recycling advice"
	Symbol for "MANUFACTURER"		
	Symbol for "THIS EQUIPMENT IS A MEDICAL DEVICE"		Symbol for "SINGLE PATIENT - MULTIPLE USE"
	Symbol for "UNIQUE DEVICE IDENTIFIER"		Symbol for "TYPE BF APPLIED PART"
	Symbol for "SERIAL NUMBER"		Symbol for "STORAGE AND TRANSPORTATION ENVIRONMENT – HUMIDITY LIMITS"
	Symbol for "European authorized representative"		Symbol for "STORAGE AND TRANSPORTATION ENVIRONMENT – PRESSURE LIMITS"
	Symbol for "Australian representative/ sponsor"		Symbol for "STORAGE AND TRANSPORTATION ENVIRONMENT – TEMPERATURE LIMITS"
	Symbol for "Importer"		Symbol for "Radio Frequency Radiation (RF)"
			Symbol for "BATCH / LOT NUMBER"

25 Network security recommendations

The following warnings detail security measures that Aktiia users should follow to ensure appropriate protection of their personal data. Failure to comply with these warnings may lead to user personal data leakage or destruction.

Only use mobile application authorized by Aktiia. Aktiia only makes its mobile application and subsequent updates available on official app stores.

Use unique credentials (username and password) for login to your Aktiia account. Safely store your password so that no other person can access it. It is recommended to regularly update your password, at least once every 3 months.

Do not let other people login to your Aktiia account on your behalf.

Version 5.0 – 2024.09.03

MD

Aktiia Init I1
Blood pressure monitor
www.aktiia.com

CE 0123

EC REP

Veranex Germany GmbH
Landsberger Str. 302, D-80687 Munich, Germany

AU REP

Emergo Australia
201 Sussex Street, Level 20, Tower II Darling Park, Sydney, NSW 2000, Australia



Aktiia SA, Bassin 8a, 2000 Neuchâtel, Switzerland